Remarks

We have amended claim 26 to correct a typographical error and to rephrase the element of the claim the requires "direct contact" between a portion of the metal chlorite and at least one acid forming component. "Direct contact" has been rephrased as "having no interposed material," with support being found in Example 1, page 26, line 23 to page 27, line 2; Example 2, page 27, line 25 to page 28, line 2; Example 3, page 28, lines 19-24; Example 4, page 29, lines 4-9; Example 5, page 29, lines 16-20; Example 6, page 30, lines 5-9; and Example 7, page 30, lines 14-22.

We have added claim 60 specify that the membrane must comprise kraft paper (support on page 22, line 9 of the specification).

We respectfully submit that the amended claim 26, and all claims depending therefrom, define an invention that is novel and non-obvious over the Derwent English abstract No. 1997-311227 of Chinese published patent specification 1104610A, because the reference fails to teach or suggest direct contact between a metal chlorite and acid forming component. The translated Chinese abstract teaches that sodium chlorite reactant must be encapsulated by Chinese wax, stearic acid, bees wax or paraffin wax. The material interposed between the sodium chlorite and the tartaric or oxalic acid creates a barrier that interferes with the sodium chlorite/acid reaction. The production of chlorine dioxide is accordingly hindered, or even prevented in areas where the barrier cannot be breached by the reaction medium, i.e., water. Furthermore, encapsulation is a relatively costly processing step, and encapsulation of a strongly oxidizing material, such as sodium chlorite, with combustible organic materials, like waxes, is dangerous due to the potentially explosive reaction that can occur between the two materials.

In contrast, the present invention requires that at least a portion of the metal chlorite and acid forming component have <u>no</u> interposed material between them, thereby promoting chlorine dioxide production, saving costs, and avoiding hazards relating to the prior art encapsulation.

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In addition, the outstanding Office action suggests that when stearic acid is utilized as the encapsulating material in the cited reference, it would constitute an acid forming component in direct contact with the sodium chlorite as required in the claims. However, the acid forming component defined in claim 26 must be selected from one of the following materials: water soluble acids, water soluble acid salts, synthetic molecular sieves, acid ion exchange resins, acid treated clays and acid treated calcined clays. While stearic acid is an acid, it is <u>not</u> water soluble, as evidenced by the attached excerpt from *Handbook of Chemistry and Physics*, CRC Press (1974). Thus, stearic acid is not an "acid forming component" as required in the present claims.

In light of the forgoing, we respectfully submit that the claims, as amended, define a novel and non-obvious invention that fully merits patent protection. We therefore respectfully request that the entire application be allowed at an early date. If there remain any issues that the Examiner believes can be resolved by discussion, the Examiner is cordially invited to contact applicant's undersigned representative at the telephone number provided below.

This amendment is being submitted after three-months, but before four-months, from the December 18, 2002 mailing date of the outstanding Office action. We hereby request a one-month extension for response, and authorization is hereby granted to charge deposit Account No. 05-1070 for the associated fee. If any other fee is required in association with this response, we hereby request and authorize the charging of the fee to deposit Account No. 05-1070.

Respectfully submitted,

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Version with markings to show changes made

Please amend claim 26 as follows:

26. (three-times amended) A device capable of producing an aqueous solution of chlorine dioxide when said device is placed into water, the device comprising a water-permeable membrane defining at least in part an enclosed space containing a mixture of at least one metal chlorite and at least one acid forming component, at least a [of] portion of said at least one metal chlorite and at least one acid forming component having no interposed material [being in direct contact], said acid forming component being selected from the group consisting of water soluble acids, water soluble acid salts, synthetic molecular sieves, acid ion exchange resins, acid treated clays and acid treated calcined clays, and wherein said metal chlorite and said acid forming component are such that they will react with each other in the presence of water but not in the substantial absence of water to produce chlorine dioxide, said membrane comprising a material which permits: (a) controlled passage of liquid water and/or water vapor into the enclosed space to thereby allow the metal chlorite and the acid forming component to react to produce chlorine dioxide and (b) passage of the so produced chlorine dioxide into a body of liquid water to produce the aqueous solution of chlorine dioxide.

Please add the following new claim 60:

60. (New) The device of claim 26, wherein the membrane comprises kraft paper.

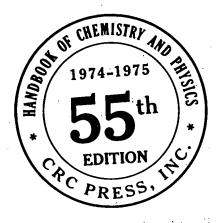


Handbook

 \mathbf{OF}

Chemistry and Physics

A Ready-Reference Book of Chemical and Physical Data



EDITOR

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In collaboration with a large number of professional chemists and physicists whose assistance is acknowledged in the list of general collaborators and in connection with the particular tables or sections involved.

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SYMBOLS AND ABBREVIATIONS

		fl	flakes	par	partial petroleum ether
[a]	specific rotation		fluorescent	peth pk	pink ³
2	slightly	fr	freezes	Ph	phenyl
>	above, more than	fr. p.	freezing point	pl	plates
<	below, less than soluble in all proportions	fum	fuming	pr pr	prisms
00	name approved by the	gel	gelatinous .	Pr	propyl
*	International Union of	gl	glacial	Prak	J. Prak. Chem.
	Chamiete (UU C.)	gold	golden	purp	purple ³
_	IR, or UV, or NMR spectra	gr	green ³	pw	powder
Ω	referenced	gran	granular gray ³	Ру	pyremidine
?	unknown	gу	hot ·	pym	pyramids
aa	acetic acid	h H	Helv. Chim. Acta	rac	racemic rectangular
abs	absolute	hex	hexagonal	rect	red
ac	acid	hp	heptane	red	resinous
Ac	acetyl	htng	heating	res rh	rhombic
ace	acetone	hx	hexane	rhd	rhombohedral
al	alcohol ²	hyd	hydrate	S	soluble
alk	alkali	hyg	hygroscopic	s	secondary ⁷
Am	J. Am. Chem. Soc.	i	insoluble	sc	scales
Am	amyl (pentyl) amorphous	i-	iso-	sec	secondary ⁷
amor	anhydrous	ign	ignites	sf	softens
anh	aqueous	in	inactive inflammable	sh	shoulder
aqu	asymmetric	inflam	infusible	silv	silvery
as atm	atmospheres	infus		sl	slightly (δ) solid
b	boiling	irid	isooctane	so	solution
B	Beilstein	iso J	I Chem. Soc.	sol solv	solvent
Ber	Chem. Ber.	joc	J. Org. Chem.	sph	sphenoidal
bipyr	n bipyramidal	L, l	levo4	spii	stable
bk	black ³	la	large	sub	sublimes
ы	blue ³	lf	leaf '	suc	supercooled
br	brown ³	lig	ligroin	sulf	sulfuric acid
bt	bright	liq	liquid	sym	symmetrical
Bu	butyl Benzene	lo	long	syr	syrup
bz C	Chem Abs	lt	light	t	tertiary ⁷
c	percentage concentration	m	melting meta-	ta	tablets
ca	about (circa)	<i>m-</i> M	molar (concentration)	tcl	triclinic tertiary ⁷
chl	chloroform	M	Merck Index, 7th Edition	tert	Tetrahedron
co	columns	mcl	monoclinic	Tet tetr	tetragonal
col	colorless	Me	methyl	TH	1 1 - 6 - 6 - 6
con	concentrated	met	metallic	to	toluene
cor	corrected	mic	r microscopic	tr	transparent
cr	crystals cyclohexane	min		trg	trigonal
сy	decomposes	mo		uno	dil undiluted
d D	line in the spectrum of	mu	normal chain, refractive	uns	
υ	sodium (subscript)	n	indev	un	
D. 0	a dextro ⁴	N	normal (concentration)	V	very vacuum
δd	slight decomposition	N	nitrogen ⁶	vac	iahla
dil	diluted	nd	needles	va: va	
dio	x dioxane	0-	ortho-	vic	F + 3 1
dis		oct	octahedral	vis	viscous
dk	dark , dl racemic⁴	og	orange ³ s ordinary organic solvents		dat volatile or volatilises
		00		vt	
dlo	MF dimethyl formamide	or		w	
E		or		w	
E. eff		or	L orthorhombic	w	
Et	ethyl	or os	- min colvents	w	.113
et	h ether ⁵	p-	_	ye	
ov.	n explodes	p-		X.	yl xyiene
ex	trap extrapolated	P	•		

No.	Name
Ωal	Abietic acid
Ωai	,methyl ester Acacetin
Qi	Acenaphthene
	a5 —,1-amino a6 —,3-amino a7 —,4-amino a8 —,5-amino
S	a9,5-bromo-
	a10,5-chloro- a11,5-iodo- ,3-nitro-
	-,1-oxo a13 5-Acenapht carboxylic
	0 a14 Acenaphthi quinone a15 3-Acenaph sulfonic a
	al6 1-Acenaph
	Ω a18 Acetaldel
	a19 —,bis(2-ethyl) a
	Ωa21 —,dieth
	0 a 2 3, 2, 4 pheny (stable, (topic) form)
	n a25 —,oxii a26 —,phe zone

Name

For I.U.C. rules of nomenclature see General Index.
 Generally means ethyl alcohol.
 The abbreviation of a color ending in "sh" is to be read as ending with the suffix "-ish," e.g., grsh means greenish.
 D. L generally mean configuration and d, l generally mean optical rotation, but there are many examples in the chemical literature for which the meaning of these symbols is ambiguous and/or interchangeable.
 Generally means diethyl ether.
 N indicates a position in the molecule.
 s and sec, or t and tert, are used as convenient.

PHYSICAL CONSTANTS OF ORGANIC COMPOUNDS (Continued)

				SICAL CON		Color.	:,	1	1	n _D			Solubility			Ref.	
	1	Name	Synonyr	ns and Formula	Mol. wt.	form. specific rotand Amer ()	ation °C	b.p.	Density	"B	w al	eth	ace	bz	solvents	 	-
1								6769	0.806740	1.452020	i		.∫ ∞	s	chl s	B13,582	1
7	Octa	cosane		₁₎₂₆ CH ₃	394.78	mcl or rh	64.5 fr 61.4	431.6 ⁷⁶⁶ 264 ¹⁰	0.77507	1.43307	٥		.\	\		B23, 1095	1
01	Octac	osane*	CH ₃ (CH ₂	1)26CH3	124.76	(bz-al) (ace or aa	1004		0.8191100	1.4313		1	\			B13, 1849	1
02	Octac	osanoic	CH ₃ (CH	₂) ₂₆ CO ₂ H	724	1	1	sub			1			, 0	chi.	B22, 459	1
	a cid		CH3(CH	₂) ₂₇ OH	410.7	1	_5	200-5 229-30	0.902240	1.46992	`	∞ °	o oc	, l "	CCI4.	1	1
			1		1	<i>c</i> 1	-,			}	1	\	1		McOF	١ هـ	1
04	9,12- cad	Octade- ienoic acid	CH,(C	acid. H₂)₄CH:CHCH₂CH	l:CII(C	pa ye or		1			11		1		1		1
	(cis	cis)*	1			232 (3.		1		.		s	s .	.	\	B22,461	1
			1		308.	275 (2.		270-5	0.8865					- \	1		1
0	se	thyl ester*	Ethyl lit	noleate. CH ₂) ₄ CH:CHCH ₂ C	H:CH((H ₂),CO ₂ C	μH,	212	1	1		- 1	-	- 1	1	1	١
Ů	1		CH3C	JH2/4CIII-		1 251 19	5 (4.0). j			1	1 1	1	}	1	1 .	D22 461	- 1
			1		1	λ ^{ε1} 233 270 (1	sh (2.1). .79)	2152	0.8886	1.463	3 ²⁰ i	5	s .	$\cdots \mid \cdot$		B22,461	- 1
	1			l linoleate.	294	40	-35	168	-70¹	`\				١	1	1	1
Ω	o6 —.	methyl ester* .	Methy	l linoleate. CH2)4CH:CHCH2C	CH:CH(CH ₂),CO ₂ ye or	col				060					B23, 1476	- 1
			1		1	λ ^{a1} 23	2 (3.78) 56-	7.5	0.868	670 1.468	9			- 1	1	Ì	- 1
	07 10	,12-Octade-	10.12-	Linoleic acid. (CH2)4CH:CHCH:(1 Z 84 CH(CH	,),CO,H	0	- 1			- }	\	1 1		. \	B23, 1068	. 1
٠.	c	adienoic acid trans, trans)*	1		1	[λ ^m 2.	33 (3.51)	167	_87 0.841	19 1.46	98¹° i	1	$ \cdots $				i
	- 1		CH.(CH2)3C:CCH2CH2	C:C(CI	1 ₂),CH ₃ 6.44			ı		\ i		.\		····	B12,772	
		11-Octa- lecadiyne*		aldehyde.	24	8.49 nd (p	eth) 55	(38–9) 261 21	2-3	1,44	.1025	. v	V	\cdots	Me	OH v Am 80, 661	3
	09 C	ctadecanal* .			ne• 3	14.56			-70 ³ ····	1	39020	ة ا	s	s	ligs	B13,565	·
	o10 -	_,dimethyl ace	tal . 1,1-I	Dimethoxyoctaucca.		1	loreth- 28	.18 31	- 610		1	į.	1	8	s chl	v B43,43	1
		Octadecane*	CH	(CH ₂) ₁₆ CH ₃		Me	OH)	cz 86 34	8.8760 0.80	518 ²⁰ 1.4	52220	i '	, ,	1	1 1	B43, 43	3
	1	,1-amino-* .	l cu	3(CH ₂)17NH ₂	• • • • •	(w)	l l	84.5	99.510			i '	v · ·	1	. 5	B43,43	
ς	012		- 1 .	3(CH ₂)17NH ₂ . CH ₃	CO₂H :	329.57 nd (al), cr (bz)	1				i }	δli	· } · ·	· i ···		
	013	,,acetate.		CU NH. HCI		305.98 orh	pl (al)	62-3		84820 1.	4631 ²⁰	• 1	v" s	s .	pe	th, B13,5	67.
	014			(3(CH ₂) ₁₇ NH ₂ . HCl		333.41 cr (al) 2	.0.2		1	.4594 ³⁰ 4531 ²⁰	; \				B13,5	66
!	2 015	,1-bromo-*		1 ₃ (CH ₂) ₁₇ Br	١.	288.95		28.6	48 ⁷⁶⁰ 0.1	3641 ²⁰ 1.	4551	1	8.	1.	ct	niv B1²,1	139
	O o16	1-chloro-*		H ₃ (CH ₂) ₁₇ Cl		١.	or if (al)	64	05-715			· \	v* }			B13,	567
		1,18-dibro		(CH ₂) ₁₈ Br		1	į	34		099420 1	.481020	i	δ	δ		B21,	626
		,1-iodo-*		H ₃ (CH ₂) ₁₇ I		380.40 lf	and ace)	}	22310			i \	s	s	.		, 2247
		L.		aicosanedioat	e	370.58		· · ·	1			i					,
		Octadecaned		C ₂ H ₃ O ₂ C(CH ₂) ₁₆ C(OH ₂) ₁₆ C(OH ₂) ₁₆ C(OH ₂) ₁₆ OH		286.50 lf	(al, AcOEt or bz), nd (bz,	97–9	210-12	,		1		1	1 1	B1'	, 1838
	02	1,18-Octa- decanediol	Į r	10(CH ₂) ₁₈ OH		1	diox)	(i) 24-6	1881-2).8475 ²⁰	1.464520	i	δ	5			
		ì	١.	-Octadecyl mercapt	an.	286.57		(ii) 28			1.429980	i	s*	v	sδ	chl. CCl ₄ , B2 ² CS ₂ , to s	
	02	1-Octa- decanethio	•	CH ₃ (CH ₂) ₁₇ SH		284.50 r	ncl lf (al) 210 (1.69)	71.5-2.0	23215			.∫ i'	δ s*	s	$\delta \delta$	chls B2	, 384
	Ωοί		ic .	Stearic acid. CH ₃ (CH ₂) ₁₆ CO ₂ H			f(al)	109	250-112			li	δ	V	sv	Circo.	2², 148
	Ωο			Stearamide. CH ₃ (CH ₂) ₁₆ CONI	-I.,	359.60	nd (al)	94	153.510			1	١.	δ	δ	peth δ	22, 360
	0	24N-pl	ienyl	Stearanilide. CH ₃ (CH ₂) ₁₆ CONI			λ ^{al} 243 (4.17)	72		0.836842	1.43628	1	1	1	1 1	chl o	6 ² , 418
		241,anhydr	ide*	Stearic anhydride. [CH ₃ (CH ₂) ₁₆ CO] ₂		1 1		28 (45–6	,\	0.907555	1.46635	° i	δ	δ	1	1 1	2 ² , 352
						374.61	pa ye	1	223	0.85520	1.4328	•	i s	1	· * · · ·	1	32, 384
	•	1		CH ₃ (CH ₂) ₁₆ CO ₂ C				27.5	21515	1	1.4523	24 .	s	٠	1	.1	
		026 ,butyl e		CH ₃ (CH ₂) ₁₆ CO ₂ (Stearyl chloride.	CH ₂) ₃ C	302.93		23	202-36	0.00023	1	ı,	, i	i ,	s	1	36 ² , 11
	Ω	o27 —.chlori	ie	CH ₃ (CH ₂) ₁₆ COC	Cl	366.64		44 (28-9		0.89023		40	, \	s i	s v	1	B2 ² , 379
		o28cyclol	exyl			```\		(i) 33.4	19910	1.0574° (0.8973	1.4349	1			1.1.	chi,	B22,353
		acter*	ster*	Ethyl stearate. CH ₃ (CH ₂) ₁₆ CO ₂	C,H,	312.54	licliath	(ii) 30 57	٠		1.4416)"°	i	i		CS25	
	12	030,hexac				508.92 -CH	lf or pl (eth, aa)		1	١.	.	060	1	δ	s* \\.		B23,10
		03U nexac	,	CH ₃ (CH ₂) ₁₆ CO		328.5	(peth)	60-1	189-91		l l	ا ``			1 1		B22,35
		0301	iroxyethyl	Glycol monostear CH ₃ (CH ₂) ₁₆ CO	rate.	нон	\"	(i) 22	5 22315	0.8498	! ° ∤				5		
		actor*	ıtyl ester .	C113(1) wax	(ii) 2	8.9 1993	1				1			1
		031 ,15001	,.	ls and abbreviations	,,,,,,,,,,			1 6000	ulas see end	of table.							